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TABLE 19-2.-F FACTORS FOR VARIOUS FUELS1

Fuel Torre	F <sub>d</sub>		F	w	F <sub>c</sub>						
Fuel Type	dscm/J	dscf/106 Btu	wscm/J	wscf/106 Btu	scm/J	scf/106 Btu					
Coal:											
Anthracite 2	2.71×10 <sup>-7</sup>	10,100	2.83×10-7	10,540	0.530×10-7	1,970					
Bituminus 2	2.63×10 <sup>-7</sup>	9,780	2.86×10 <sup>-7</sup>	10,640	0.484×10 <sup>-7</sup>	1,800					
Lignite	2.65×10 <sup>-7</sup>	9,860	3.21×10 <sup>-7</sup>	11,950	0.513×10 <sup>-7</sup>	1,910					
Oil 3	2.47×10-7	9,190	2.77×10-7	10,320	0.383×10-7	1,420					
Gas:.											
Natural	2.34×10 <sup>-7</sup>	8,710	2.85×10 <sup>-7</sup>	10,610	0.287×10 <sup>-7</sup>	1,040					
Propane	2.34×10 <sup>-7</sup>	8,710	2.74×10 <sup>-7</sup>	10,200	0.321×10 <sup>-7</sup>	1,190					
Butane	2.34×10 <sup>-7</sup>	8,710	2.79×10 <sup>-7</sup>	10,390	0.337×10 <sup>-7</sup>	1,250					
Wood	2.48×10 <sup>-7</sup>	9,240			0.492×10 <sup>-7</sup>	1,830					
Wood Bark	2.58×10 <sup>-7</sup>	9,600			0.516×10 <sup>-7</sup>	1,920					
Municipal	2.57×10 <sup>-7</sup>	9,570			0.488×10 <sup>-7</sup>	1,820					
Solid Waste											

 $<sup>^1</sup>$  Determined at standard conditions: 20 °C (68 °F) and 760 mm Hg (29.92 in Hg)  $^2$  As classified according to ASTM D 388.  $^3$  Crude, residual, or distillate.

TABLE 19-3.—VALUES FOR To 95\*

n¹	t <sub>0.95</sub>	n¹	t <sub>0.95</sub>	n¹	t <sub>0.95</sub>
2	6.31	8	1.89	22–26	1.71
3	2.42	9	1.86	27-31	1.70
4	2.35	10	1.83	32-51	1.68
5	2.13	11	1.81	52-91	1.67
6	2.02	12-16	1.77	92-151	1.66
7	1.94	17–21	1.73	152 or more	1.65

<sup>&</sup>lt;sup>1</sup>The values of this table are corrected for n-1 degrees of freedom. Use n equal to the number (H) of hourly average data points

METHOD 20—DETERMINATION OF NITROGEN OX-IDES, SULFUR DIOXIDE, AND DILUENT EMIS-SIONS FROM STATIONARY GAS TURBINES

## 1. Principle and Applicability

1.1 Applicability. This method is applicable for the determination of nitrogen oxides  $(NO_x)$ , sulfur dioxide  $(SO_2)$ , and a diluent gas, either oxygen (O2) or carbon dioxide (CO2), emissions from stationary gas turbines. For the NOx and diluent concentration determinations, this method includes: (1) Measurement system design criteria; (2) Analyzer performance specifications and performance test procedures; and (3) Procedures for emission testing.

1.2 Principle. A gas sample is continuously extracted from the exhaust stream of a stationary gas turbine; a portion of the sample stream is conveyed to instrumental analyzers for determination of NO<sub>x</sub> and diluent content. During each  $NO_{\mathrm{x}}$  and diluent determination, a separate measurement of SO2 emissions is made, using Method 6, or its equivalent. The diluent determination is used to adjust the NOx and SO2 concentrations to a reference condition.

## 2. Definitions

2.1 Measurement System. The total equipment required for the determination of a gas concentration or a gas emission rate. The system consists of the following major subsystems:

2.1.1 Sample Interface. That portion of a system that is used for one or more of the following: sample acquisition, sample transportation, sample conditioning, or protection of the analyzers from the effects of the stack effluent.

2.1.2  $NO_x$  Analyzer. That portion of the system that senses  $NO_x$  and generates an output proportional to the gas concentration

2.1.3 O2 Analyzer. That portion of the system that senses O2 and generates an output proportional to the gas concentration.

2.1.4 CO<sub>2</sub> Analyzer. That portion of the system that senses CO2 and generates an output proportional to the gas concentration.

2.1.5 Data Recorder. That portion of the measurement system that provides a permanent record of the analyzer(s) output. The data recorder may include automatic data reduction capabilities.

2.2 Span Value. The upper limit of a gas concentration measurement range that is specified for affected source categories in the applicable part of the regulations.

2.3 Calibration Gas. A known concentration of a gas in an appropriate diluent gas.

2.4 Calibration Error. The difference between the gas concentration indicated by the measurement system and the known concentration of the calibration gas.